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(54) Title: LOW RESIDUE AQUEOUS HARD SURFACE CLEANING COMPOSITIONS PARTICULARLY ADAPTED FOR CLEANING GLASS AND GLOSSY HARD SURFACES

(57) Abstract: Aqueous based cleaning and disinfecting compositions particularly useful for cleaning glass and polished hard surfaces comprise at least one linear 6-18C alkyl sulfate surfactant, at least one glycoside-based surfactant, a solvent system containing an alkylene glycol ether and 1-6C alcohol, and water.

LOW RESIDUE AQUEOUS HARD SURFACE CLEANING COMPOSITIONS PARTICULARLY ADAPTED FOR CLEANING GLASS AND GLOSSY HARD SURFACES

Cleaning compositions are commercially important products and enjoy a wide field of utility in assisting in the removal of dirt and grime from surfaces, especially glass and glossy hard surfaces (i.e., glazed ceramic tiles, polished metals, enameled metal surfaces, glazed porcelain). While the art is replete with various formulations which provide some cleaning benefit and perhaps some disinfecting benefit to surfaces, there is a real and continuing need for such further formulations.

Thus, it is among the objects of the invention to provide improved aqueous cleaning compositions which are especially useful in cleaning, especially hard surfaces particularly glass and other glossy hard surfaces. Such a composition is particularly useful for use "as-is" by the ultimate user. It is a further object of the invention to provide a process for cleaning hard surfaces, which process comprises the step of: providing an aqueous cleaning composition as outlined herein, and applying an effective amount of the same to a surface, especially a hard surface requiring such cleaning treatment.

These and other objects of the invention shall be more apparent from a reading of the specification and of the claims attached.

According to one aspect of the present invention there is provided an aqueous cleaning composition which provides disinfecting and cleaning characteristics to treated surfaces, particularly hard surfaces, which comprises the following constituents:

- (A) at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant, preferably predominantly a linear lauryl sulfate;
- (B) at least one surfactant based on a glycoside;
 - (C) a solvent system containing an alkylene glycol ether solvent, especially propylene glycol n-butyl ether, ethylene glycol hexyl ether, further with a C₁-C₆ alcohol, especially where the C₁-C₆ alcohol is isopropanol;
 - (D) water.

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The compositions may include one or more further optional additive constituents, sometimes referred to as adjuvants, in minor, but effective amounts. By way of non-limiting example, such optional additives include: coloring agents such as dyes and pigments, fragrances, other pH adjusting agents, pH buffer compositions, chelating agents, rheology modification agents as well as one or more further surfactant compounds, in particular nonionic, amphoteric or zwitterionic surfactant compounds. Desirably, in order to reduce the likelihood of undesired buildup upon treated surfaces, especially hard surfaces, the amounts of these additive constituents are present in only minor amounts, i.e., less than 10%, preferable less than 5% wt. based on the total weight of the aqueous cleaning composition being provided herein.

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The inventive compositions include (A) at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant. Such materials are commercially available as anionic surfactants and are frequently provided in a salt form in an aqueous carrier. Typically such salt forms include counterions based on alkali or alkaline earth metals. It is known that such sulfate surfactants may be provided as a technical grade mixture which may include sulfates which include various C₆-C₁₈ alkyl chains. According to the present invention it is preferred that a preponderance of alkyl groups be C₁₂ groups, viz., be linear C₁₂ alkyl sulfates. Still more preferably the consist of at least 70%, more preferably at least 85%, still more preferably comprise at least 95% of the linear alkyl groups be C₁₂ alkyl groups. Such materials are commercially available, *inter alia*, under the trade name Sulfotex® WAQ-LCX (ex. Henkel Corp.) of which various technical grade aqueous mixtures are presently commercially available.

The linear C_6 - C_{18} alkyl sulfate surfactants comprise 0 - 10%wt., yet more preferably 0.01-5%wt. but most preferably comprise 0.01 - 3%wt. of the compositions of which they form a part.

The compositions according to the present invention also include (B) at least one surfactant based on a glycoside. Exemplary suitable compounds include alkyl monoglycosides and polyglycosides are prepared generally by reacting a monosaccharide, or a compound hydrolyzable to a monosaccharide with an alcohol such as a fatty alcohol in an acid medium.

Exemplary alkyl glycoside surfactants suitable for use in the practice of this invention may be represented by formula I below:

$$RO$$
— $(R_1O)_V$ – $(G)_XZ_b$

wherein:

R is a monovalent organic radical containing from about 6 to about 30, preferably from about 8 to about 18 carbon atoms;

R₁ is a divalent hydrocarbon radical containing from about 2 to about 4 carbon atoms, especially ethyl and propyl radicals;

O is an oxygen atom;

y is a number which has an average value from about 0 to about 1 and is preferably 0;

G is a moiety derived from a reducing saccharide containing 5 or 6 carbon atoms; and

x is a number having an average value from about 1 to 5 (preferably from 1.1 to 2);

Z is selected O₂M¹,

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O(CH₂), CO₂M¹, OSO₃M¹, or O(CH₂)SO₃M¹; R₂ is (CH₂)CO₂M¹ or CH=CHCO₂M¹; with the proviso that Z can be O₂M¹ only if Z is in place of a primary hydroxyl group in which the primary hydroxylbearing carbon atom,

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-CH2OH, is oxidized to form a

group);

b is a number of from 0 to 3x+1 preferably an average of from 0.5 to 2 per glycosyl group;

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M¹ is H⁺ or an organic or inorganic cation, such as, for example, an alkali metal, ammonium, monoethanolamine, or calcium.

As defined in Formula I above, R is generally the residue of a fatty alcohol having from about 8 to 30 and preferably 8 to 18 carbon atoms.

Most preferably, the inventive compositions include an alkylpolyglycoside compound according to the structure:

wherein:

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R is an alkyl group, preferably a linear alkyl chain, which comprises C_8 to C_{16} alkyl groups;

x is an integer value of from 0-3, inclusive.

Examples of such alkylpolyglycoside compounds according to this structure include: where R is comprised substantially of C₈ and C₁₀ alkyl chains yielding an average value of about 9.1 alkyl carbons per molecule (GLUCOPON 220 UP, GLUCOPON 225 DK); where R is comprised of C₈, C₁₀, C₁₂, C₁₄ and C₁₆ alkyl carbons yielding an average value of about 10.3 alkyl carbons per molecule (GLUCOPON 425); where R is comprised substantially of C₁₂, C₁₄ and C₁₆ alkyl carbons yielding an average value of about 12.8 alkyl carbons per molecule (GLUCOPON 600 UP, GLUCOPON 625 CSUP, and GLUCOPON 625 FE, all of which are available from Henkel Corp., Ambler PA.) Also useful as the alkylpolyglycoside compound is TRITON CG-110 (Union Carbide Corp.).

Further examples of commercially available alkylglycosides as described above include, for example, GLUCOPON 325N which is described as being a 50% C₉-C₁₁ alkyl polyglycoside, also commonly referred to as D-glucopyranoside (from Henkel Corp, Ambler PA). Particularly preferred as the alkylpolyglycoside compounds are those illustrated in the Examples. These alkylpolyglycosides may be present in any amount. It will be understood that this such a minimum amount will vary widely, and is in part dependent upon the molecular weight of the alkylpolyglycoside utilized in a formulation, but desirably at least about 0.01%wt. should be present. More preferably the polyglycoside comprises from 0.02%wt. to 10%wt. of the compositions of which it forms a part. Further and particularly

preferred examples of alkylpolyglycosides are described with reference to the Examples.

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The compositions of the invention include (C) a solvent system containing an alkylene glycol ether solvent, further with a C1-C6 alcohol, especially where the C1-C6 alcohol is isopropanol. Particularly useful alkylene glycol ethers include C3-C20 glycol ethers. Specific illustrative examples of useful alkylene glycol ether solvents include: propylene glycol methyl ether, dipropylene glycol methyl ether, tripropylene glycol methyl ether, propylene glycol isobutyl ether, ethylene glycol methyl ether, ethylene glycol ethyl ether, ethylene glycol butyl ether, diethylene glycol phenyl ether, propylene glycol phenol ether, and mixtures thereof. More preferably employed as the solvent is one or more of the group consisting of ethylene glycol n-butyl ether, diethylene glycol n-butyl ether, and mixtures thereof. Preferably the (C) solvent system includes both propylene glycol n-butyl ether with ethylene glycol hexyl ether, and in certain especially preferred embodiments propylene glycol n-butyl ether and ethylene glycol hexyl ether are the sole glycol ethers of the (C) solvent system. Propylene glycol n-butyl ether is known to the art. It is commercially available as Dowanol® PnB (ex. Dow Chem. Co., Midland, MI). The propylene glycol n-butyl ether may be present in amounts of from 0.01%wt. - 6.0 wt., however is advantageously present in amounts of from 0.01 - 4%wt. Ethylene glycol hexyl ether is also, per se, known to the art. It is commercially available as Hexyl Cellosolve® (ex. Union Carbide Corp.). The ethylene glycol hexyl ether may be present in amounts of from 0.01%wt. - 5.0 %wt., however is advantageously present in amounts of from 0.01 - 2%wt.

The inventors have observed that the compositions according to the present invention exhibit desirable evaporation and drying properties (e.g., relatively uniform drying with no streaking, no mottling). Certain particularly preferred embodiments of the inventive compositions form a substantially uniform film during evaporative drying subsequent to application on a hard surface.

The compositions of the invention also include a C_1 - C_6 alcohol as part of the (C) solvent system. Such include for example methanol, ethanol, n-propanol, isopropanol as well as the various positional isomers of butanol, pentanol and hexanol. The inclusion of such alcohols have been found by the present inventor to

even further improve in the evaporation of the inventive composition in a relatively even manner such that it tends to form a relatively uniform film layer during the drying process. This effect has been generally described above in conjunction with glycol n-butyl ether and ethylene glycol hexyl ether. A further benefit of the inclusion of such alcohols is in the solvency which they may provide to certain stains as well. Of these, the inclusion of isopropanol is most preferred. The C₁-C₆ alcohol may be present in amounts of from 0.01%wt. - 10%wt., however is advantageously present in amounts of from 0.01 - 60%wt.

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According to certain particularly preferred embodiments, the (C) a solvent system consists solely of propylene glycol n-butyl ether, ethylene glycol hexyl ether and isopropanol to the exclusion of other C_1 - C_6 alcohols and other glycol ethers.

As is noted above, the compositions according to the invention are aqueous in nature, and include (D) as a further essential constituent. Water is added in order to provide to 100% by weight of the compositions of the invention, and is preferably deionized water.

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Certain preferred embodiments of compositions according to the invention may be categorized as "non-food contact surface sanitizing" compositions as they exhibit antimicrobial efficacy against at least *Staphylococcus aureus*, *Salmonella cholerasuis*, and *Enterobacter aerogenes* in accordance with the "ASTM Standard Test Method for Efficacy of Sanitizers Recommended for Inanimate Non-Food Contact Surfaces, E 1153 - 87", or broad spectrum disinfecting efficacy against at least *Staphylococcus aureus*, or *Salmonella cholerasuis* (preferably against both) by the "AOAC Official Method 961.02 Germicidal Spray Products as Disinfectants" (per AOAC Official Methods of Analysis, 16th Edition (1995)). Both of these tests are known to those skilled in the art.

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As noted, the compositions may include one or more optional additives which by way of non-limiting example include: coloring agents such as dyes and pigments, fragrances and fragrance solubilizers, pH adjusting agents, pH buffering agents, chelating agents, rheology modification agents, as well as one or more further nonionic surfactant compounds. Desirably, in order to reduce the likelihood of undesired buildup upon treated surfaces, especially hard surfaces, the total amounts of such optional additives is less than about 10% wt. but are desirably significantly less,

such as less than about 2.5%wt. based on the total weight of the aqueous cleaning and disinfecting composition being provided herein. Optimally, the amounts of such further optional additives is kept to a minimum in order to minimize the amounts of non-volatile constituents in the compositions as a whole, which tend to contribute to an undesired streaky or mottled appearance of the composition during drying.

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Useful as chelating agents include those known to the art, including by way of non-limiting example; aminopolycarboxylic acids and salts thereof wherein the amino nitrogen has attached thereto two or more substituent groups. Preferred chelating agents include acids and salts, especially the sodium and potassium salts of ethylenediaminetetraacetic acid, diethylenetriaminepentaacetic acid, N-hydroxyethylethylenediaminetriacetic acid, and of which the sodium salts of ethylenediaminetetraacetic acid may be particularly advantageously used. Such chelating agents may be omitted, or they may be included in generally minor amounts such as from 0 - 0.5 %wt. based on the weight of the chelating agents and/or salt forms thereof. Desirably, such chelating agents are included in the present inventive composition in amounts from 0 - 0.5%wt., but are most desirably present in reduced weight percentages from about 0 - 0.2%wt.

The compositions according to the invention optionally but desirably include an amount of a pH adjusting agent or pH buffer composition. Such compositions include many which are known to the art and which are conventionally used. By way of non-limiting example pH adjusting agents include phosphorus containing compounds, monovalent and polyvalent salts such as of silicates, carbonates, and borates, certain acids and bases, tartrates and certain acetates. Further exemplary pH adjusting agents include mineral acids, basic compositions, and organic acids, which are typically required in only minor amounts. By way of further non-limiting example pH buffering compositions include the alkali metal phosphates, polyphosphates, pyrophosphates, triphosphates, tetraphosphates, silicates, metasilicates, polysilicates, carbonates, bicarbonates, hydroxides, and mixtures of the same. Certain salts, such as the alkaline earth phosphates, carbonates, hydroxides, can also function as buffers. It may also be suitable to use as buffers such materials as aluminosilicates (zeolites), borates, aluminates and certain organic materials such as gluconates, succinates, maleates, and their alkali metal salts. These may be included in any effective amount

which may be used to adjust and maintain the pH of the compositions of the invention to the desired pH range, which is usually in the range of from about 4 - 12. These should be screened however to ensure that they do not undesirably deposit residues upon the surfaces being treated. Particularly useful are sodium hydroxide, and sodium bicarbonate, which are present in certain particularly preferred embodiments of the invention.

Further optional, but advantageously included constituents are one or more coloring agents which find use in modifying the appearance of the compositions and enhance their appearance from the perspective of a consumer or other end user. Known coloring agents may be incorporated in the compositions in any effective amount to improve or impart to compositions a desired appearance or color. Such a coloring agent or coloring agents may be added in a conventional fashion, i.e., admixing to a composition or blending with other constituents used to form a composition.

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Further optional, but desirable constituent include fragrances, natural or synthetically produced. Such fragrances may be added in any conventional manner, admixing to a composition or blending with other constituents used to form a composition, in amounts which are found to be useful to enhance or impart the desired. scent characteristic to the composition, and/or to cleaning compositions formed therefrom. In addition to a fragrance, it is frequently desirable to include a fragrance solubilizer which assists in the dispersion, solution or mixing of the fragrance constituent in an aqueous base. These include known art compounds, including condensates of 2 to 30 moles of ethylene oxide with sorbitan mono- and tri-C₁₀-C₂₀ alkanoic acid known to be useful as nonionic surfactants. Further examples of such suitable surfactants include water soluble nonionic surfactants of which many are commercially known and by way of non-limiting example include the primary aliphatic alcohol ethoxylates, secondary aliphatic alcohol ethoxylates, alkylphenol ethoxylates and ethylene-oxide-propylene oxide condensates on primary alkanols, and condensates of ethylene oxide with sorbitan fatty acid esters.. This fragrance solubilizer component is added in minor amounts, particularly amount which are found effective in aiding in the solubilization of the fragrance component, but not in any significantly greater proportion, such that it would be considered as a detergent

constituent. Such minor amounts recited herein are generally up to about 0.3% by weight of the total composition but is more generally an amount of about 0.1% by weight and less, and preferably is present in amounts of about 0.05% by weight and less.

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As an optional constituent, the compositions may include one or more nonionic surfactant compounds in amounts which are effective in improving the overall cleaning efficacy of the compositions being taught herein, while at the same time in amounts which do not undesirably diminish the germicidal efficacy of the inventive compositions or which undesirably increase the likelihood to form or deposit surface residues onto the treated surfaces. Such nonionic surfactant compounds are known to the art.. Practically any hydrophobic compound having a carboxy, hydroxy, amido, or amino group with a free hydrogen attached to the nitrogen can be condensed with ethylene oxide or with the polyhydration product thereof, polyethylene glycol, to form a water soluble nonionic surfactant compound. Further, the length of the polyethylenoxy hydrophobic and hydrophilic elements may be varied. Exemplary nonionic compounds include the polyoxyethylene ethers of alkyl aromatic hydroxy compounds, e.g., alkylated polyoxyethylene phenols, polyoxyethylene ethers of long chain aliphatic alcohols, the polyoxyethylene ethers of hydrophobic propylene oxide polymers, and the higher alkyl amine oxides.

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To be mentioned as particularly useful nonionic surfactants are alkoxylated linear primary and secondary alcohols such as those commercially available under the tradenames PolyTergent® SL series (Olin Chemical Co., Stamford CT), Neodol® series (Shell Chemical Co., Houston TX); as alkoxylated alkyl phenols including those commercially available under the tradename Triton® X series (Union Carbide Chem. Co., Danbury CT).

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Further useful optional surfactants include amine oxides, such as lauryl dimethyl amine oxides, betaines and sarcosinate based surfactants.

Such constituents as described above as essential and/or optional constituents include known art compositions, include those described in *McCutcheon's Emulsifiers* and *Detergents (Vol.1)*, *McCutcheon's Functional Materials (Vol. 2)*, North American Edition, 1991; *Kirk-Othmer, Encyclopedia of Chemical Technology*, 3rd Ed., Vol. 22, pp. 346-387, the contents of which are herein incorporated by reference.

According to one particularly preferred aspect of the present invention there is provided an aqueous cleaning composition which provides both disinfecting and cleaning characteristics to treated surfaces, particularly hard surfaces, which comprises the following constituents:

- 5 (A) 0.01 10%wt. of at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant, preferably predominantly a linear lauryl sulfate;
 - (B) 0.01 10%wt. of at least one surfactant based on a glycoside;
 - a solvent system containing: 0.01 11%wt. of one or more alkylene glycol ether solvents, (especially 0.01 6.0%wt. propylene glycol n-butyl ether, 0.1 5%wt. ethylene glycol hexyl ether) with 0.1 10%wt. of a C₁-C₆ alcohol, especially where the C₁-C₆ alcohol is isopropanol;
 - (D) to 100%wt., water.
 and further, 0.01 2%wt. of a pH buffer or pH adjusting agent, especially one or more hydroxides, carbonates or bicarbonates,
 and further 0 10%wt. of optional additives as described herein.

The compositions of the invention can be prepared in a conventional manner such as by simply mixing the constituents in order to form the ultimate aqueous cleaning composition. The order of addition is not critical.

The compositions according to the invention are useful in the cleaning and/or sanitizing of surfaces, especially hard surfaces, having deposited soil thereon. The compositions are particularly effective in the removal of oleophilic soils (viz., oily soils) particularly of the type which are typically encountered in kitchens and other food preparation environments. In such a process, cleaning and disinfecting of such surfaces comprises the step of applying a soil releasing and disinfecting effective amount of a composition as taught herein to such a soiled surface. Afterwards, the compositions are optionally but desirably wiped, scrubbed or otherwise physically contacted with the hard surface, and further optionally, may be subsequently rinsed from such a cleaned and disinfected hard surface.

The hard surface cleaner composition provided according to the invention can be desirably provided as a ready to use product in a manually operated spray dispensing container and is thus ideally suited for use in a consumer "spray and wipe" application. In such an application, the consumer generally applies an effective

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amount of the cleaning composition using the pump and within a few moments thereafter, wipes off the treated area with a rag, towel, or sponge, usually a disposable paper towel or sponge. To ensure effective sanitization or disinfection, a longer contact time, generally of 10 minutes is required.

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In a yet a further embodiment, the compositions according to the invention may be formulated so that they may be useful in conjunction with an "aerosol" type product wherein they are discharged from a pressurized aerosol container. If the inventive compositions are used in an aerosol type product, it is preferred that corrosion resistant aerosol containers such as coated or lined aerosol containers be used. Such are preferred as they are known to be resistant to the effects of acidic formulations. Known art propellants such as liquid propellants as well as propellants of the non-liquid form, i.e., pressurized gases, including carbon dioxide, air, nitrogen, hydrocarbons as well as others may be used. Also, while satisfactory for use, fluorocarbons may be used as a propellant but for environmental and regulatory reasons their use is preferably avoided.

The composition of the present invention, whether as described herein or in a concentrate or super concentrate form, can also be applied to a hard surface by using a wet wipe. The wipe can be of a woven or non-woven nature. Fabric substrates can include nonwoven or woven pouches, sponges, in the form of abrasive or nonabrasive cleaning pads. Such fabrics are known commercially in this field, and are often referred to as wipes. Such substrates can be resin bonded, hydroentanged, thermally bonded, meltblown, needlepunched or any combination of the former.

The nonwoven fabrics may be a combination of wood pulp fibers and textile length synthetic fibers formed by well known dry-form or wet-lay processes. Synthetic fibers such as rayon, nylon, orlon and polyester as well as blends thereof can be employed. The wood pulp fibers should comprise about 30 to about 60 percent by weight of the nonwoven fabric, preferably about 55 to about 60 percent by weight, the remainder being synthetic fibers. The wood pulp fibers provide for absorbency, abrasion and soil retention whereas the synthetic fibers provide for substrate strength and resiliency.

The substrate of the wipe may also be a film forming material such as a water soluble polymer. Such self-supporting film substrates may be sandwiched between

layers of fabric substrates and heat sealed to form a useful substrate. The free standing films can be extruded utilizing standard equipment to devolatilize the blend. Casting technology can be used to form and dry films, or a liquid blend can be saturated into a carrier and then dried in a variety of known methods.

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The compositions of the present invention are absorbed onto the wipe to form a saturated wipe. The wipe can then be sealed individually in a pouch which can then be opened when needed or a multitude of wipes can be placed in a container for use on an as-needed basis. The container, when closed, is sufficiently sealed to prevent evaporation of any components from the compositions.

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Whereas the present invention is intended to be produced and provided in the "ready-to-use" form described above, nothing in this specification shall be understood as to limit the use of the composition according to the invention with a further amount of water to form a cleaning solution therefrom. The aqueous compositions according to the invention may be used, and are preferably used "as-is" without further dilution, they may also be used with a further aqueous dilution. Such dilutions include ratios (w%/w%, or v%/v%) of composition:water concentrations of from 1:0, to extremely dilute dilutions such as 1:10,000. Desirably however, in order to ensure disinfection the compositions should be used "as is", that is to say without further dilution.

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The following examples illustrate the superior properties of the formulations of the invention and particular preferred embodiments of the inventive compositions. The terms "parts by weight" or "percentage weight" are used interchangeably in the specification and in the following Examples wherein the weight percentages of each of the individual constituents are indicated in weight percent based on the total weight of the composition of which it forms a part, unless indicated otherwise.

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Examples:

Exemplary formulations illustrating certain preferred embodiments of the inventive compositions and described in more detail in Table 1 below were formulated generally in accordance with the following protocol.

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Into a suitably sized vessel, a measured amount of water was provided after which the constituents were added in the following sequence: surfactants, solvents followed by the remaining constituents, including any optional constituents. All of the

constituents were supplied at room temperature, and mixing of the constituents was achieved by the use of a mechanical stirrer with a small diameter propeller at the end of its rotating shaft. Mixing, which generally lasted from 5 minutes to 120 minutes was maintained until the particular exemplary formulation appeared to be homogeneous. The exemplary compositions were readily pourable, and retained well mixed characteristics (i.e., stable mixtures) upon standing. It is to be noted that the constituents might be added in any order, but it is preferred that water be the initial constituent provided to a mixing vessel or apparatus as it is the major constituent and addition of the further constituents thereto is convenient. The exact compositions of the example formulations are listed on Table 1, below wherein are indicated the weight percentages of the individual constituents, based on a total weight of 100% weight.

Table 1	
	Ex.1
sodium lauryl sulfate	0.40
alkyl polyglucoside	0.15
propylene glycol n-butyl ether	1.5
ethylene glycol hexyl ether	1.0
isopropanol	3.5
sodium hydroxide	0.041
sodium bicarbonate	0.05
fragrance	0.05
di water	to 100

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The amounts indicated on Table 1 relating to each constituent indicate the "actives weight" of the identified constituent. The specific identity and source of the particular constituents recited in Table 1 are also disclosed in Table 2 below.

Table 2	
sodium lauryl sulfate	Sulfotex® WAQ-LCX, 30%wt. actives, from Henkel
alkyl polyglucoside	APG 325N, 50%wt. actives, from Henkel
propylene glycol n-butyl ether	Dowanol® PnB, 100%wt. actives, ex. Dow Chemical Co.
ethylene glycol hexyl ether	Hexyl Cellosolve, 100%wt. actives, ex. Union Carbide Corp.
isopropanol	isopropanol, 100%wt. actives, ex. Aldrich Chem. Co.
sodium hydroxide	anhydrous, 25%wt. actives, ex. Aldrich Chem. Co.
sodium bicarbonate	anhydrous, 100%wt. actives, ex. Aldrich Chem. Co.
fragrance	proprietary composition
di water	deionized water

The compositions of Table 1 were evaluated in accordance with one or more of the further tests described below.

Evaluation of Antimicrobial Efficacy:

The exemplary formulation described on Table 1 above was evaluated in order to evaluate its antimicrobial efficacy against *Staphylococcus aureus* (gram positive type pathogenic bacteria) (ATCC 6538), and *Enterobacter aerogenes aureus* (gram negative type pathogenic bacteria) (ATCC 13048) in accordance with the "ASTM Test Standard Test Method for Efficacy of Sanitizers Recommended for Inanimate Non-Food Contact Surfaces, E 1153-87). As is appreciated by the skilled practitioner in the art, the results of this test indicates log reduction of test organisms which are subjected to a test composition based on Ex.1. The results of the antimicrobial testing are indicated on Table 3, below.

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Tabl 3		
Enterobacter aerog	genes	
Formula:	Log ₁₀ recovery	Log ₁₀ reduction
Ex.1	0	5.46
Control	5.46	n/a
Staphylococcus au	reus	
Formula:	Log ₁₀ recovery	Log ₁₀ reduction
Ex.1	0	5.89
Control	5.89	n/a

As a control, ("Control" in Table 3) an aqueous composition containing 0.01%wt. of an ethoxylated phenolic surfactant (Triton-X100®, ex. Union Carbide) was also tested. As may be seen from the results indicated above, the compositions according to the invention provide excellent sanitization of hard surfaces, while the compositions based on the ethoxylated phenolic surfactant performed poorly. It is to be noted that a Log₁₀ reduction of 3 or more indicates a passing sanitization score in the test.

The formulation described on Table 1 was also evaluated in order to evaluate its antimicrobial efficacy against two series of 60 test samples of *Staphylococcus aureus* (gram positive type pathogenic bacteria) (ATCC 6538), and a single series of 30 test organisms of), *Salmonella choleraesuis* (gram negative type pathogenic bacteria) (ATCC 10708), in accordance with the protocol described within the "AOAC Official Method 961.02 Germicidal Spray Products as Disinfectants" (per AOAC Official Methods of Analysis, 16th Edition (1995)). The results of the antimicrobial testing are indicated on Table 4, below. The reported results indicate the number of test slides with live test organisms/number of test slides at the conclusion of the test for the example formulation and organism tested, as well as the recovery counts.

Table 4				
	live test organisms/number of test slides			
Ex.1, Staphylococcus aureus, (1st test)	0/60			
Ex.1, Staphylococcus aureus (2 nd test)	0/60			
Ex.1, Salmonella choleraesuis	0/30			

Evaluation of Cleaning Efficacy:

The compositions according the invention are expected to provide good cleaning.

5 Evaluation of Evaporation and Drying Characteristics:

The compositions of the invention are expected to have good evaporative and satisfactory drying characteristics.

Claims:

1. A low residue aqueous cleaning and disinfecting composition which comprises:

- (A) at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant
- (B) at least one surfactant based on a glycoside;
- (C) a solvent system containing an alkylene glycol ether solvent, further with a C_1 - C_6 alcohol;
- (D) water.

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- 2. The low residue aqueous hard surface cleaning and disinfecting composition according to claim 1 wherein the surfactant (A) is predominantly a linear lauryl sulfate.
- The low residue aqueous hard surface cleaning and disinfecting composition according to claim 1 wherein the sole alkylene glycol ether solvents are propylene glycol n-butyl ether, and ethylene glycol hexyl ether.
 - 4. The low residue aqueous hard surface cleaning and disinfecting composition according to claim 1 wherein the solvent system consists solely of propylene glycol n-butyl ether, ethylene glycol hexyl ether and a C₁-C₆ alcohol.
 - 5. The low residue aqueous hard surface cleaning and disinfecting composition according to claim 1 wherein the solvent system consists solely of propylene glycol n-butyl ether, ethylene glycol hexyl ether and isopropanol.
 - A composition according to claim 1 further characterized in that the composition forms a substantially uniform film during evaporative drying subsequent to application on a hard surface.

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7. A composition according to claim 1 which further comprises a pH adjusting agent or a pH buffering agent.

8. A composition according to claim 1 comprising per 100%wt.:

- (A) 0.01 10%wt. of at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant;
- (B) 0.01 10%wt. of at least one surfactant based on a glycoside;
 - (C) a solvent system containing: 0.01 11%wt. of one or more alkylene glycol ether solvents, and 0.1 10%wt. of a C_1 - C_6 alcohol;
 - (D) to 100%wt., water.

 a pH buffer or pH adjusting agent;
- and further 0 10%wt. of optional additives as described herein.
 - 9. A composition according to claim 1 comprising per 100%wt.:
 - (A) 0.01 10%wt. of at least one anionic a linear C₆-C₁₈ alkyl sulfate surfactant;
 - (B) 0.01 10%wt. of at least one surfactant based on a glycoside;
 - (C) a solvent system containing: 0.01 6.0%wt. propylene glycol n-butyl ether, 0.1 5%wt. ethylene glycol hexyl ether and 0.1 10%wt. of a C_1 - C_6 alcohol;
 - (D) to 100%wt., water.a pH buffer or pH adjusting agent;and further 0 10%wt. of optional additives as described herein.
 - 10. The composition according to claim 8 wherein the compositions are characterized by forming a substantially uniform film during evaporative drying after being applied to a hard surface.
 - 11. The composition according to claim 9 wherein the compositions are characterized by forming a substantially uniform film during evaporative drying after being applied to a hard surface

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12. A process for the cleaning and disinfecting of a hard surface in need of such treatment which comprises the step of:
applying an effective amount of the composition according to claim 1.

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13. A low residue aqueous cleaning and disinfecting composition substantially as described with reference to the example.

INTERNATIONAL SEARCH REPORT

Interr nal Application No PCT/GB 00/01859

A. CLASSIFICATION OF SUBJECT MATTER
1PC 7 C11D3/48 C11D1/83 C11D3/20 A01N25/30 A01N25/02 //C11D1/66,C11D1/14 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) IPC 7 C11D A01N Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) WPI Data, EPO-Internal, PAJ C. DOCUMENTS CONSIDERED TO BE RELEVANT Category ' Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. WO 98 50510 A (UNILEVER) 1,2,4-13 Α 12 November 1998 (1998-11-12) page 8, line 9 - line 15; examples 34-37 WO 98 21305 A (RECKITT & COLMAN INC) Α 1.3 - 1322 May 1998 (1998-05-22) abstract page 3, line 1 - line 15; claims 1-6 EP 0 875 550 A (HENKEL KGAA) 1,2,4-6, A 4 November 1998 (1998-11-04) 8-13 page 3, line 2; claims 1,9; examples Further documents are listed in the continuation of box C. Patent family members are listed in annex. Special categories of cited documents : "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance invention "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention filing date cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled in the art. other means "P" document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 5 September 2000 12/09/2000 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Saunders, T Fax: (+31-70) 340-3016

INTERNATIONAL SEARCH REPORT

information on patent family members

Intern nal Application No PCT/GB 00/01859

Patent document cited in search report		Publication date	Patent family member(s)			Publication date
WO 98505	10 A	A 12-11-1998		7651698 0980419		27-11-1998 23-02-2000
WO 98213	05 A	22-05-1998	GB	2319179	Α	20-05-1998
			AU	4341897	Α	03-06-1998
			BR	9713005	Α	25-01-2000
			CN	1237203	Α	01-12-1999
			EP	0938542	Α	01-09-1999
			US	5891392	A	06-04-1999
EP 08755	50 A	04-11-1998	DE	19717950	A	05-11-1998